| L Number | Hits | Search Text | DB | Time stamp |
|----------|------|---|-----------|------------------|
| 1 | 0 | surfactancy near15 antigen near15 binding | USPAT; | 2003/09/05 14:30 |
| | | | US-PGPUB; | |
| | | | EPO; JPO; | |
| | | | DERWENT | |
| 2 | 1 | surfactancy near15 antigen | USPAT; | 2003/09/05 14:30 |
| | | | US-PGPUB; | 1 |
| | | | EPO; JPO; | 1 |
| | | | DERWENT | |
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| | | | US-PGPUB; | |
| | | | EPO; JPO; | |
| | | * | DERWENT |] |
| 4 | 3 | surfactancy near10 increase | USPAT; | 2003/09/05 14:31 |
| | | | US-PGPUB; | |
| | | | EPO; JPO; | |
| | | | DERWENT | |

surfactancy(P)(solid or bead or particle or particulate)(P)(binding or affinity)

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L5
            0 FILE CERAB
L6
            0 FILE METADEX
L7
            O FILE USPATFULL
TOTAL FOR ALL FILES
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=> surfactancy (10A) bead
            1 FILE CAPLUS
L9
            0 FILE BIOTECHNO
L10
L11
            O FILE COMPENDEX
L12
            0 FILE ANABSTR
L13
            0 FILE CERAB
            O FILE METADEX
L14
L15
            1 FILE USPATFULL
TOTAL FOR ALL FILES
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=> d 19 ibib abs total
    ANSWER 1 OF 1 CAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER:
                    2001:886639 CAPLUS
DOCUMENT NUMBER:
                        136:17677
TITLE:
                       No wash bead assay, kit and procedure
INVENTOR(S):
                       Hechinger, Mark K.
PATENT ASSIGNEE(S):
                        Cytometry Applications, Inc., USA
SOURCE:
                        PCT Int. Appl., 28 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT:
PATENT INFORMATION:
    PATENT NO.
                    KIND DATE
                                         APPLICATION NO. DATE
                    A1 20011206 WO 2001-US40837 20010604
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    WO 2001092887
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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                                     US 2001-873866 20010604
EP 2001-939955 20010604
    US 2002004199 A1 20020110
    EP 1292829
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PRIORITY APPLN. INFO.:
                                       US 2000-209437P P 20000602
                                       WO 2001-US40837 W 20010604
    A method of making a no wash bead based assay comprises prepg. a first
AΒ
    reagent comprising a buffer, and prepg. a second reagent comprising a
    than 5 are prepd., including washing the beads in the buffer to form a
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AB A method of making a no wash bead based assay comprises prepg. a first reagent comprising a buffer, and prepg. a second reagent comprising a protein. Beads of preselected size and having a coeff. of variation less than 5 are prepd., including washing the beads in the buffer to form a bead-buffer matrix and reducing the surfactancy of the beads to an effective amt. Thereafter, an antigen for detecting the presence of a target species is added to the bead-buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixt. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixt. and thereafter the mixt. is incubated. The second reagent is added to the bead-antigen mixt. to reduce or eliminate non-specific binding sites.

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=> surfactancy (15A) reduce (20A) bead L17 0 FILE CAPLUS L18 0 FILE BIOTECHNO L19 0 FILE COMPENDEX L20 0 FILE ANABSTR 0 FILE CERAB L21 0 FILE METADEX L22L23 O FILE USPATFULL

TOTAL FOR ALL FILES

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1 FILE CAPLUS

PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'BEAD) (P) REDUCE'

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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'BEAD) (P) REDUCE'

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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'BEAD) (P) REDUCE'

L30 O FILE METADEX L31 1 FILE USPATFULL

TOTAL FOR ALL FILES

2 (SURFACTANCY(10A) BEAD)(P) REDUCE

=> d l32 ibib abs total

L32 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:886639 CAPLUS

DOCUMENT NUMBER: 136:17677

INVENTOR(S): No wash bead assay, kit and procedure

Hechinger, Mark K. Cytometry Applications, Inc., USA PATENT ASSIGNEE(S):

SOURCE: PCT Int. Appl., 28 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

| PAT | PENT | NO. | | KI | ND : | DATE | | | A. | PPLI | CATI | ON NO | ο. | DATE | | | |
|---------------|------------|-------------|-----|-------------------------|------|-----------------|-----|-----|------|------|----------|-------|------|------|-----|-----|-----|
| | | | | | | | | | - | | | | | | | | |
| WO 2001092887 | | A1 20011206 | | | | WO 2001-US40837 | | | | | 20010604 | | | | | | |
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| | | LU, | LV, | MA, | MD, | MG, | MK, | MN, | MW, | MX, | MZ, | NO, | NZ, | PL, | PT, | RO, | RU, |
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| ΕP | EP 1292829 | | A: | 1 : | 2003 | 0319 | | E | P 20 | 01-9 | 3995 | 5 | 2001 | 0604 | | | |
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IE, SI, LT, LV, FI, RO, MK, CY, AL, TR

PRIORITY APPLN. INFO.:

US 2000-209437P P 20000602 WO 2001-US40837 W 20010604

AB A method of making a no wash bead based assay comprises prepg. a first reagent comprising a buffer, and prepg. a second reagent comprising a protein. Beads of preselected size and having a coeff. of variation less than 5 are prepd., including washing the beads in the buffer to form a bead-buffer matrix and reducing the surfactancy of the beads to an effective amt. Thereafter, an antigen for detecting the presence of a target species is added to the bead-buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixt. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixt. and thereafter the mixt. is incubated. The second reagent is added to the bead-antigen mixt. to reduce or eliminate non-specific binding sites.

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L32 ANSWER 2 OF 2 USPATFULL on STN

ACCESSION NUMBER: 2002:8201 USPATFULL

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K., Pasadena, CA, UNITED STATES

NUMBER DATE

PRIORITY INFORMATION: US 2000-209437P 20000602 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: COLIN P ABRAHAMS, 5850 CANOGA AVENUE, SUITE 400,

WOODLAND HILLS, CA, 91367

NUMBER OF CLAIMS: 26 EXEMPLARY CLAIM: 1 LINE COUNT: 793

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A method of making a no wash bead based assay comprises preparing a first reagent comprising a buffer, and preparing a second reagent comprising a protein. Beads of preselected size and having a coefficient of variation less than 5% are prepared, including washing the beads in the buffer to form a bead-buffer matrix and reducing the surfactancy of the beads to an effective amount. Thereafter, an antigen for detecting the presence of a target species is added to the bead-buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixture. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixture and thereafter the mixture is incubated. The second reagent is added to the bead-antigen mixture to reduce or eliminate non-specific binding sites.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

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^{=&}gt; file .meeting

^{&#}x27;EVENTLINE' IS NOT A VALID FILE NAME

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=> surfactancy(P)(solid or bead or particle or particulate)(P)binding
L33
             0 FILE AGRICOLA
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FIELD CODE - 'AND' OPERATOR ASSUMED 'URFACTANCY(P) (SOLID'
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FIELD CODE - 'AND' OPERATOR ASSUMED 'RTICULATE) (P) BINDING'
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L40
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=> surfactancy(P)(solid or bead or particle or particulate)(P)(affinity or binding)
             0 FILE AGRICOLA
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stable mean droplet size of about 300 nm. The stability of the mucin-emulsified systems was also evaluated by measuring turbidity

changes with time, which allowed a comparison with similar emulsions stabilized by the Pluronic.RTM. surfactants in the same concentration. Thus, mucin showed its ability to establish more stable and more efficient oil-water emulsion systems. Since mucin is a glycoprotein, and hence biodegradable, our results suggest that mucin might serve as an ideal biological surfactant for the stabilization of emulsion systems intended for biomedical and pharmaceutical applications.

=> file .jacob

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=> surfactancy(P)(solid or bead or particle or particulate)(P)(binding or affinity)

L51 1 FILE CAPLUS L52 0 FILE BIOSIS L53 0 FILE MEDLINE 0 FILE EMBASE L54 L55 2 FILE USPATFULL

TOTAL FOR ALL FILES

L56 3 SURFACTANCY(P)(SOLID OR BEAD OR PARTICLE OR PARTICULATE)(P)(BIND ING OR AFFINITY)

=> dup rem

ENTER L# LIST OR (END):156 PROCESSING COMPLETED FOR L56

L57 3 DUP REM L56 (0 DUPLICATES REMOVED)

=> d 157 ibib abs total

L57 ANSWER 1 OF 3 USPATFULL on STN

ACCESSION NUMBER: 2003:30295 USPATFULL

TITLE:

Particles with improved solubilization capacity

INVENTOR(S):

Anderson, David, Colonial Heights, VA, UNITED STATES

NUMBER KIND DATE -----US 2003022242 A1 20030130 US 2002-176112 A1 20020621 (10) PATENT INFORMATION: APPLICATION INFO.:

NUMBER DATE

PRIORITY INFORMATION: US 2001-300476P 20010623 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: WHITHAM, CURTIS & CHRISTOFFERSON, P.C., 11491 SUNSET

HILLS ROAD, SUITE 340, RESTON, VA, 20190

NUMBER OF CLAIMS: 204 EXEMPLARY CLAIM: 1

NUMBER OF DRAWINGS: 1 Drawing Page(s)

LINE COUNT: 3885

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A particle is disclosed that comprises a first volume of hydrophobe-rich material with tunable dissolution and solubilization characteristics and a distinct second volume of nanostructured nonlamellar liquid crystalline material, said second volume containing said first domain and being capable of being in equilibrium with said first volume. Preferably, the nanostructured nonlamellar liquid crystalline material is capable of being in equilibrium with a polar solvent or a water-immiscible solvent or both.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L57 ANSWER 2 OF 3 USPATFULL on STN

ACCESSION NUMBER: 2002:8201 USPATFULL

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K., Pasadena, CA, UNITED STATES

NUMBER DATE

PRIORITY INFORMATION: US 2000-209437P 20000602 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: COLIN P ABRAHAMS, 5850 CANOGA AVENUE, SUITE 400,

WOODLAND HILLS, CA, 91367

NUMBER OF CLAIMS: 26
EXEMPLARY CLAIM: 1
LINE COUNT: 793

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

Amethod of making a no wash bead based assay comprises preparing a first reagent comprising a buffer, and preparing a second reagent comprising a protein. Beads of preselected size and having a coefficient of variation less than 5% are prepared, including washing the beads in the buffer to form a bead -buffer matrix and reducing the surfactancy of the beads to an effective amount. Thereafter, an antigen for detecting the presence of a target species is added to the bead -buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixture. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixture and thereafter the mixture is incubated. The second reagent is added to the bead-antigen mixture to reduce or eliminate non-specific binding sites.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L57 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:886639 CAPLUS

DOCUMENT NUMBER: 136:17677

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K.

PATENT ASSIGNEE(S):

Cytometry Applications, Inc., USA

SOURCE:

PCT Int. Appl., 28 pp. CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

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    WO 2001092887 A1 20011206 WO 2001-US40837 20010604
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             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRIORITY APPLN. INFO.:
                                       US 2000-209437P P 20000602
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A method of making a no wash bead based assay comprises prepg. a first reagent comprising a buffer, and prepg. a second reagent comprising a protein. Beads of preselected size and having a coeff. of variation less than 5 are prepd., including washing the beads in the buffer to form a bead-buffer matrix and reducing the surfactancy of the beads to an effective amt. Thereafter, an antigen for detecting the presence of a target species is added to the bead-buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixt. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixt. and thereafter the mixt. is incubated. The second reagent is added to the bead-antigen mixt. to reduce or eliminate non-specific binding sites.

REFERENCE COUNT:

5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

WO 2001-US40837 W 20010604

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CA INDEXING COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)
=> surfactancy(P)(solid or bead or particle or particulate)(P)(binding or affinity)
L58
             1 FILE CAPLUS
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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'URFACTANCY(P) (SOLID' PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'RTICULATE) (P) (BINDING' 1 FILE BIOTECHNO PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'URFACTANCY(P) (SOLID' PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH FIELD CODE - 'AND' OPERATOR ASSUMED 'RTICULATE) (P) (BINDING' L60 0 FILE COMPENDEX L61 0 FILE ANABSTR L62 0 FILE CERAB

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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH

FIELD CODE - 'AND' OPERATOR ASSUMED 'RTICULATE) (P) (BINDING'

L63 1 FILE METADEX L64 2 FILE USPATFULL

TOTAL FOR ALL FILES

5 SURFACTANCY(P) (SOLID OR BEAD OR PARTICLE OR PARTICULATE)(P)(BIND ING OR AFFINITY)

=> dup rem ENTER L# LIST OR (END):165 PROCESSING COMPLETED FOR L65 L66 5 DUP REM L65 (0 DUPLICATES REMOVED)

=> d 166 ibib abs total

L66 ANSWER 1 OF 5 USPATFULL on STN

ACCESSION NUMBER: 2003:30295 USPATFULL

TITLE: Particles with improved solubilization capacity

INVENTOR(S): Anderson, David, Colonial Heights, VA, UNITED STATES

NUMBER KIND DATE -----PATENT INFORMATION: US 2003022242 A1 20030130 APPLICATION INFO.: US 2002-176112 A1 20020621 (10)

NUMBER DATE -----

PRIORITY INFORMATION: US 2001-300476P 20010623 (60) Utility

DOCUMENT TYPE: FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: WHITHAM, CURTIS & CHRISTOFFERSON, P.C., 11491 SUNSET

HILLS ROAD, SUITE 340, RESTON, VA, 20190

NUMBER OF CLAIMS: 204

EAEMPLARY CLAIM: 1
NUMBER OF DRAWINGS: 1 Drawing Page(s)
LINE COUNT: 2007

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A particle is disclosed that comprises a first volume of hydrophobe-rich AB material with tunable dissolution and solubilization characteristics and a distinct second volume of nanostructured nonlamellar liquid crystalline material, said second volume containing said first domain and being capable of being in equilibrium with said first volume. Preferably, the nanostructured nonlamellar liquid crystalline material is capable of being in equilibrium with a polar solvent or a water-immiscible solvent or both.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L66 ANSWER 2 OF 5 USPATFULL on STN

ACCESSION NUMBER: 2002:8201 USPATFULL

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K., Pasadena, CA, UNITED STATES

NUMBER KIND DATE US 2002004199 A1 20020110 US 2001-873866 A1 20010604 (9) PATENT INFORMATION: APPLICATION INFO.:

> NUMBER DATE -----

PRIORITY INFORMATION: US 2000-209437P 20000602 (60)

DOCUMENT TYPE: Utility FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: COLIN P ABRAHAMS, 5850 CANOGA AVENUE, SUITE 400,

WOODLAND HILLS, CA, 91367

NUMBER OF CLAIMS: EXEMPLARY CLAIM: LINE COUNT: 793

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method of making a no wash bead based assay comprises preparing a first reagent comprising a buffer, and preparing a second reagent comprising a protein. Beads of preselected size and having a coefficient of variation less than 5% are prepared, including washing the beads in the buffer to form a bead -buffer matrix and reducing the surfactancy of the beads to an effective amount. Thereafter, an antigen for detecting the presence of a target species is added to the bead -buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixture. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixture and thereafter the mixture is incubated. The second reagent is added to the bead-antigen mixture to reduce or eliminate non-specific binding sites.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L66 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2003 ACS on STN

ACCESSION NUMBER: 2001:886639 CAPLUS

DOCUMENT NUMBER: 136:17677

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K.

PATENT ASSIGNEE(S): Cytometry Applications, Inc., USA

SOURCE: PCT Int. Appl., 28 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

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PATENT NO.
                  KIND DATE
                                         APPLICATION NO. DATE
     WO 2001092887 Al 20011206 WO 2001-US40837 20010604
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            HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
            LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,
            SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU,
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            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                     US 2001-873866 20010604
EP 2001-939955 20010604
     US 2002004199
                     A1 20020110
     EP 1292829
                      A1
                           20030319
            AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
             IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRIORITY APPLN. INFO.:
                                       US 2000-209437P P 20000602
                                       WO 2001-US40837 W 20010604
    A method of making a no wash bead based assay comprises prepg. a
     first reagent comprising a buffer, and prepg. a second reagent comprising
     a protein. Beads of preselected size and having a coeff. of
     variation less than 5 are prepd., including washing the beads in
     the buffer to form a bead-buffer matrix and reducing the
     surfactancy of the beads to an effective amt.
     Thereafter, an antigen for detecting the presence of a target species is
     added to the bead-buffer matrix such that the antigen attaches
     to the beads to form a bead-antigen mixt. The
     surfactancy of the beads facilitates attachment of the
     antigen thereto. Buffer is added to the bead-antigen mixt. and
     thereafter the mixt. is incubated. The second reagent is added to the
    bead-antigen mixt. to reduce or eliminate non-specific
    binding sites.
REFERENCE COUNT:
                        5
                              THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS
                              RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
L66
     ANSWER 4 OF 5 BIOTECHNO COPYRIGHT 2003 Elsevier Science B.V. on STN
ACCESSION NUMBER:
                        1999:29457809
                                       BIOTECHNO
TITLE:
                        Effects of mucin addition on the stability of
                        oil-water emulsions
AUTHOR:
                        Shi L.; Miller C.; Caldwell K.D.; Valint P.
CORPORATE SOURCE:
                        K.D. Caldwell, Center for Biopolymers at Interfaces,
                        University of Utah, Contact Lens Division, Salt Lake
                        City, UT 84112, United States.
                        E-mail: karin.caldwell@biosurf.uu.se
SOURCE:
                        Colloids and Surfaces B: Biointerfaces, (1999), 15/3-4
                        (303-312), 18 reference(s)
                        CODEN: CSBBEQ ISSN: 0927-7765
PUBLISHER ITEM IDENT.:
                        S092777659900096X
DOCUMENT TYPE:
                        Journal; Article
COUNTRY:
                        Netherlands
LANGUAGE:
                        English
SUMMARY LANGUAGE:
                        English
AN
     1999:29457809
                     BIOTECHNO
     In this work, bovine submaxillary gland mucin (BSM) was used as an
AB
     emulsifier to stabilize oil-water emulsion systems. Prior to use,
     commercial BSM was purified by jacalin affinity chromatography.
     Emulsions consisting of 5% mineral oil in phosphate buffered saline (PBS)
     were prepared through the addition of different amounts of purified mucin
     followed by sonication using either of two methods: (1) low energy input
     for a long time (2 h), or (2) high energy input for a short time (20 s).
     The surfactancy property of mucin was investigated by surface
     tension measurements, which showed the BSM to greatly reduce the surface
```

tension of PBS. Compared to several synthetic surfactants of the Pluronic.RTM. type, mucin showed comparable or better surface activity than F68, F88 and F108 products in dilute solutions. The formed emulsions had a mean droplet size that decreased monotonically with increasing concentration of mucin until a plateau was reached at concentrations around 0.1% by weight. The stability of these emulsions was evaluated by monitoring their average droplet size during a 33-day period. Emulsions with more than 0.25% mucin showed a constant mean size throughout the period. Specifically, an emulsion produced with 0.95% mucin showed a stable mean droplet size of about 300 nm. The stability of the mucin-emulsified systems was also evaluated by measuring turbidity changes with time, which allowed a comparison with similar emulsions stabilized by the Pluronic.RTM. surfactants in the same concentration. Thus, mucin showed its ability to establish more stable and more efficient oil-water emulsion systems. Since mucin is a glycoprotein, and hence biodegradable, our results suggest that mucin might serve as an ideal biological surfactant for the stabilization of emulsion systems intended for biomedical and pharmaceutical applications.

L66 ANSWER 5 OF 5 METADEX COPYRIGHT 2003 CSA on STN

ACCESSION NUMBER:

1996(1):12-103 METADEX

TITLE:
AUTHOR:

Carbonaceous inclusions in aluminum alloys. Quintero-Sayago, O. (Universidad Simon Bolivar);

Ramirez, S.C. (Universidad Simon Bolivar)

SOURCE:

Georgia Institute of Technology School of Materials Science and Engineering. Atlanta, GA 30332-0385, USA. 1994. 66-73, Photomicrographs, Diffraction Patterns,

16 ref.

Conference: The 4th International Conference on Aluminum Alloys, Their Physical and Mechanical

Properties. Vol. I, Atlanta, GA, USA, 11-16 Sept. 1994

Conference Article

COUNTRY:

DOCUMENT TYPE:

United States

LANGUAGE:

English

AB Chemical distribution, taken inside two carbon containing inclusions of idiomorphic and dendritic morphologies, is used as a proof to discuss their probable dormation mechanism in Al melt. The nucleation potential of the Al-Si-C particles in the liquid is explained, based on the interactions between silicon and carbon, partially depending on the hypothesis of qualitative semi-empirical concepts of alloy theory when applied to the liquid state. Even though the tendency for Al-Si bonding in liquid is more favorable to exist, high surfactancy on halide salts promote wetting. This enhances the generation of carbonaceous inclusions in the melt.

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=> surfactancy(P)(solid or bead or particle or particulate)
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PROXIMITY OPERATOR LEVEL NOT CONSISTENT WITH
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L71
             0 FILE CERAB
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            1 FILE METADEX
L73
            19 FILE USPATFULL
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TOTAL FOR ALL FILES

L74 23 SURFACTANCY (P) (SOLID OR BEAD OR PARTICLE OR PARTICULATE)

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L87
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L88
L89
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TOTAL FOR ALL FILES
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L90 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2003 ACS on STN
ACCESSION NUMBER: 2001:886639 CAPLUS
DOCUMENT NUMBER:
                       136:17677
TITLE:
                        No wash bead assay, kit and procedure
INVENTOR(S):
                       Hechinger, Mark K.
PATENT ASSIGNEE(S):
                        Cytometry Applications, Inc., USA
SOURCE:
                        PCT Int. Appl., 28 pp.
                        CODEN: PIXXD2
DOCUMENT TYPE:
                        Patent
LANGUAGE:
                        English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:
    PATENT NO. KIND DATE
                                   APPLICATION NO. DATE
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    WO 2001092887 A1 20011206 WO 2001-US40837 20010604
        W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
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            HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT,
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            SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU,
            ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
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            BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
    US 2002004199 A1 20020110 US 2001-873866 20010604 EP 1292829 A1 20030319 EP 2001-939955 20010604
        R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
            IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
PRIORITY APPLN. INFO.:
                                       US 2000-209437P P 20000602
                                       WO 2001-US40837 W 20010604
    A method of making a no wash bead based assay comprises prepg. a
AR
     first reagent comprising a buffer, and prepg. a second reagent comprising
     a protein. Beads of preselected size and having a coeff. of
     variation less than 5 are prepd., including washing the beads in
     the buffer to form a bead-buffer matrix and reducing the
     surfactancy of the beads to an effective amt.
     Thereafter, an antigen for detecting the presence of a target species is
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=> dup rem

added to the bead-buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixt. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead -antigen mixt. and thereafter the mixt. is incubated. The second reagent is added to the bead-antigen mixt. to reduce or eliminate

REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L90 ANSWER 2 OF 4 USPATFULL on STN

non-specific binding sites.

ACCESSION NUMBER: 2003:112721 USPATFULL

TITLE: Self assembling monolayer compositions

INVENTOR(S): Guire, Patrick E., Eden Prairie, MN, UNITED STATES Taton, Kristin S., Little Canada, MN, UNITED STATES

NUMBER KIND DATE -----PATENT INFORMATION: APPLICATION INFO.: US 2003077452 A1 20030424 US 2002-163012 A1 20020604 (10)

RELATED APPLN. INFO.: Continuation of Ser. No. US 2001-907303, filed on 17

Jul 2001, GRANTED, Pat. No. US 6444318

DOCUMENT TYPE: Utility APPLICATION FILE SEGMENT:

LEGAL REPRESENTATIVE: FREDRIKSON & BYRON, P.A., 1100 International Centre,

900 Second Avenue South, Minneapolis, MN, 55402

NUMBER OF CLAIMS: EXEMPLARY CLAIM: LINE COUNT: 1618

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A surface coating composition for providing a self-assembling monolayer, in stable form, on a material surface or at a suitable interface, as well as a method of preparing such a composition and a method of using such a composition to coat a surface, such as the surface of an implantable medical device, in order to provide the surface with desirable properties. The method provides the covalent attachment of a SAM to a surface in a manner that substantially retains or improves the characteristics and/or performance of both the SAM and the surface itself. Covalent attachment is accomplished by the use of one or more latent reactive groups, e.g., provided by either the surface and/or by the SAM-forming molecules themselves.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L90 ANSWER 3 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2002:224347 USPATFULL

Self assembling monolayer compositions TITLE:

Guire, Patrick E., Eden Prairie, MN, United States INVENTOR(S):

Taton, Kristin S., Little Canada, MN, United States

PATENT ASSIGNEE(S): Surmodics, Inc., Eden Prairie, MN, United States (U.S.

corporation)

NUMBER KIND DATE -----US 6444318 B1 20020903 US 2001-907303 20010717 PATENT INFORMATION: 20010717 (9) APPLICATION INFO.:

DOCUMENT TYPE: Utility FILE SEGMENT: GRANTED

PRIMARY EXAMINER: Boykin, Terressa M. LEGAL REPRESENTATIVE: Fredrikson & Byron, P.A.

NUMBER OF CLAIMS: 28 EXEMPLARY CLAIM:

NUMBER OF DRAWINGS: 11 Drawing Figure(s); 9 Drawing Page(s)

LINE COUNT:

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB A surface coating composition for providing a self-assembling monolayer, in stable form, on a material surface or at a suitable interface, as well as a method of preparing such a composition and a method of using such a composition to coat a surface, such as the surface of an implantable medical device, in order to provide the surface with desirable properties. The method provides the covalent attachment of a SAM to a surface in a manner that substantially retains or improves the characteristics and/or performance of both the SAM and the surface itself. Covalent attachment is accomplished by the use of one or more latent reactive groups, e.g., provided by either the surface and/or by the SAM-forming molecules themselves.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

L90 ANSWER 4 OF 4 USPATFULL on STN

ACCESSION NUMBER: 2002:8201 USPATFULL

TITLE: No wash bead assay, kit and procedure

INVENTOR(S): Hechinger, Mark K., Pasadena, CA, UNITED STATES

NUMBER DATE

PRIORITY INFORMATION: US 2000-209437P 20000602 (60)

DOCUMENT TYPE: Utility
FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: COLIN P ABRAHAMS, 5850 CANOGA AVENUE, SUITE 400,

WOODLAND HILLS, CA, 91367

NUMBER OF CLAIMS: 26 EXEMPLARY CLAIM: 1 LINE COUNT: 793

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

A method of making a no wash bead based assay comprises preparing a first reagent comprising a buffer, and preparing a second reagent comprising a protein. Beads of preselected size and having a coefficient of variation less than 5% are prepared, including washing the beads in the buffer to form a bead -buffer matrix and reducing the surfactancy of the beads to an effective amount. Thereafter, an antigen for detecting the presence of a target species is added to the bead -buffer matrix such that the antigen attaches to the beads to form a bead-antigen mixture. The surfactancy of the beads facilitates attachment of the antigen thereto. Buffer is added to the bead-antigen mixture and thereafter the mixture is incubated. The second reagent is added to the bead-antigen mixture to reduce or eliminate non-specific binding sites.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.